Docket No. 20784/4-CON

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

APPLICANT:

Demuth et al.

ART UNIT: Not yet assigned

Date: December 11, 2003

SERIAL NO.:

Not yet assigned

FILED:

December 11, 2003

FOR: PRODRUGS OF DP IV-INHIBITORS

CERTIFICATE OF EXPRESS MAILING

I hereby certify that this document (along with any paper referred to as being attached or enclosed) is being deposited with the United States Postal Service "Express Mail Post Office to Addressee" service under 37 CFR 1.10, Express Mail No: EV 264411154 US on the date shown below and is addressed to: MAIL STOP PATENT APPLICATION, Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

Sandra J. Wittrup

INFORMATION DISCLOSURE STATEMENT SUBMITTING REFERENCES ON COMPACT DISC

MAIL STOP PATENT APPLICATION Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

Pursuant to Applicant(s) duty of disclosure, the information listed in the attached form PTO-1449 is brought to the attention of the Examiner.

The citation of the listed items is not a representation that they constitute a complete or exhaustive listing of the relevant art or that the references are prior art. The items listed are submitted in good faith, but are not intended to substitute for the Examiner's search. It is hoped, however, that in addition to apprising the Examiner of these particular items, they will assist in identifying fields of search and in making as full and complete a search as possible.

The filing of this information disclosure statement is not an admission that the information cited herein is, or is considered to be, material to patentability as defined in 37 C.F.R. § 1.56(b).

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In accordance with the requirement under 37 C.F.R. 1.98 (3)(i), the following are concise explanations of the relevance, as presently understood, with regard to those items submitted herewith that are not in the English language:

I. LIST OF PATENTS, PUBLICATIONS OR OTHER INFORMATION

The patents, publications or other information submitted for consideration by the Office are listed on PTO-1449, attached hereto.

II. COPIES

a. Submitted herewith on Compact Disc is a legible copy of (i) each U.S and foreign patent; (ii) each publication or that portion which caused it to be listed; and (iii) all other information or that portion which caused it to be listed.

III. CONCISE EXPLANATION OF THE RELEVANCE

(check at least one box)

- a. Except as may be indicated below in (b), all of the patents, publications or other information are in the English language or were cited in an English language Search Report, a copy of which is attached hereto (concise explanation not required).
- b. A concise explanation of the relevance of all patents, publications or other information listed that is not in the English language is as follows:

FRENCH LANGUAGE FR 2 696 740 Applicants have relied on an English language abstract in determining that this patent apparently relates to Dimethylbiguanide drug derivatives and their medical applications.

FRENCH LANGUAGE FR 2 085 665 Applicants have relied on an English language abstract in determining that this patent apparently relates to biguanide substitutes possessing a hypoglycemic property.

GERMAN LANGUAGE DT 25 42 598 A1 Applicants have relied on an English language abstract in determining that this patent apparently relates to biguanide salts as well as a method for preparing such salts. In addition to this the invention apparently covers pharmaceutical compounds which contain such salts.

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GERMAN LANGUAGE WO 97/40832 Applicants have relied on an English language abstract in determining that this patent apparently relates to the use of a method for reducing in the blood of a mammal by administration of effectors the enzyme activity of dipeptidyl peptidase (DP IV) or enzyme activity similar to DP IV, the endogenous (or additionally exogenously administered) insulinotropic peptide gastric inhibitory polypeptide 1-42 and glucagon-like peptide amide-1 7-36 are decomposed in a causal sequence to a reduced extent by DP IV enzymes or those similar to DP IV.

JAPANESE PATENT JP 4-334357A2: Applicants have relied on an English language abstract in determining that this patent ACYL DERIVATIVE HAVING ENZYME-INHIBITING ACTION, apparently relates to a compound having a prolyl endopeptibase activity-inhibiting action and useful as an antidement agent, especially an anti-amnestic agent.

GERMAN LANGUAGE DE 299 09 210 U1 Applicants have relied on an English language abstract in determining that this patent apparently relates to dipeptide compounds or compounds analogous to dipeptide compounds, which are made of an amino acid and a thiazolidine or pyrrolidine group, and to their salts. The invention further relates to the use of these compounds in the treatment of impaired glucose tolerance, glucosuria, hyperlipidemia, metabolic acidosis, diabetes mellitus, diabetic neuropathy and nephropathy as well as secondary diseases of diabetes mellitus in mammals.

GERMAN LANGUAGE DE 198 26 972 A1 Applicants have relied on an English language abstract in determining that this patent apparently relates to the use of inhibitors (I) of dipeptidyl peptidase IV (DP IV) or analogous enzymes for inhibiting the activation and proliferation of keratinocytes (including malignantly transformed keratinocytes) and tumor cells derived from keratinocytes.

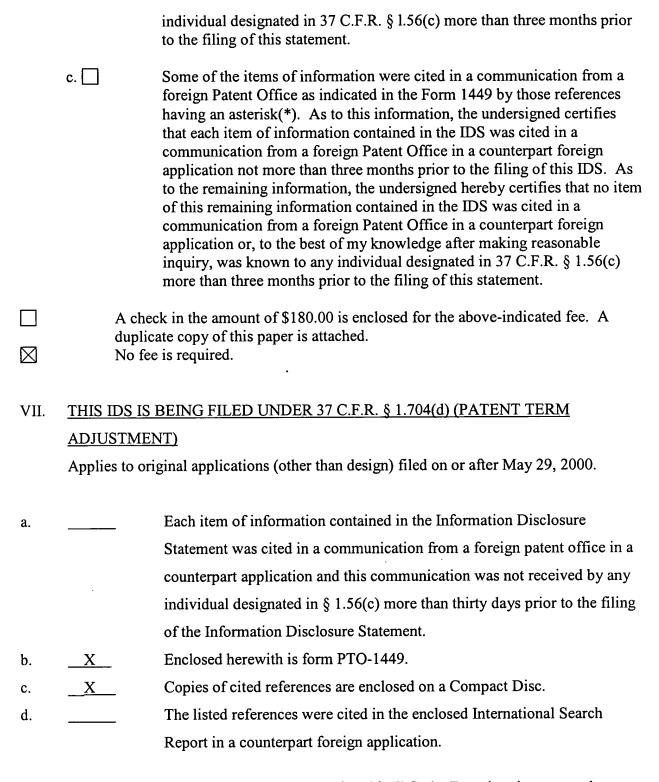
GERMAN LANGUAGE WO 99/61431 Applicants have relied on an English language abstract in determining that this patent apparently relates to dipeptide compounds or compounds analogous to dipeptide compounds, which are made of an amino acid and a thiazolidine or pyrrolidine group, and to their salts. The invention further relates to the use of these compounds in the treatment of impaired glucose tolerance, glucosuria, hyperlipidemia, metabolic acidosis, diabetes mellitus, diabetic neuropathy and nephropathy as well as secondary diseases of diabetes mellitus in mammals.

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	c	The following additional information is provided for the Examiner's consideration:							
	<u>FEES</u>								
IV.	THIS IDS IS	BEING FILED UNDER 37 C.F.R. § 1.97(b) x)							
	a. 🔀	with the application or within three months of the filing date of a national application (37 C.F.R. § 1.97(b) (1)). No fee or certification is required.							
	b. 🗌	within three months of the date of entry of the national stage as set forth in §1.491 in an international application (37 C.F.R. § 1.97(b) (2)). No fee or certification is required.							
	c. 🗌	before the mailing date of a first Action on the merits (37 C.F.R. § 1.97(b) (3)). No fee or certification is required. In the event that a first Office Action on the merits has been issued, please consider this IDS under 37 C.F.R. § 1.97(c) and see the certification under 37 C.F.R. § 1.97(e)below, or, if no certification has been made, charge our deposit account a fee in the amount of \$180.00 as required by 37 C.F.R. § 1.17(p).							
V.	THIS IDS IS (check one bo	BEING FILED UNDER 37 C.F.R. § 1.97(c):							
	1.97(c) (1)) or	iling date of a Final Office Action under 37 C.F.R. § 1.113 (See 37 C.F.R. § r before the mailing date of a Notice of Allowance under 37 C.F.R. § 1.311 c. § 1.97(c) (2)).							
	a. No cer	rtification; therefore, a fee in the amount of \$180.00 is required by 37 C.F.R. § 1.17(p).							
	b. See th	or e certification below. No fee is required.							
VI.	CERTIFICAT	TION UNDER 37 C.F.R. § 1.97(e) (check only one box)							
The ur	The undersigned hereby certifies that								
	a. 🗌	each item of information contained in the IDS was cited in a communication from a foreign Patent Office in a counterpart foreign application not more than three months prior to the filing of this IDS; or							
	b. 🗌	no item of information contained in the IDS was cited in a communication from a foreign Patent Office in a counterpart foreign application or, to the best of my knowledge after making reasonable inquiry, was known to any							

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If the Examiner has any questions concerning this IDS, the Examiner is requested to contact the undersigned. If it is determined that this IDS has been filed under the wrong rule, the

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PTO is requested to consider this IDS under the proper rule (with a petition, if necessary) and charge any additional fees to Deposit Account No. 50-0369.

Date: December 11, 2003

Respectfully submitted,

John C. Serio, Reg. No. 39,023

BROWN RUDNICK BERLACK ISRAELS LLP

Attorney(s) for Applicants Customer No. 21710

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FORM PTO-1449 INFORMATION DISCLOSURE STATEMENT			ATTY. DOCKET NO. 20784/4-CON SERIAL NO. Not yet assigned				
				APPLICANT(S): Demuth et al.	-		
				FILING DATE: December 11, 2003		RT UNIT: yet assign	ed
			UNITED STATE	ES PATENT DOCUMENTS			
EXAM. INITIAL	1 1		DATE	INVENTOR	CLASS	SUB CLASS	FIL. DATE IF APPR
	AA	2,961,377	11/22/1960	Shapiro et al.	167	65	
	AB	3,174,901	03/23/1965	Sterne	167	65	
	AC	3,879,541	04/22/1975	Kabbe et al.	424	326	
	AD	3,960,949	06/01/1976	Ahrens et al.	260	564 B	
	AE 4,028,402 06/07/1977 AF 4,935,493 06/19/1990 AG 5,433,955 07/18/1995 AH 5,462,928 10/31/1995 AI 5,512,549 04/30/1996	Fischer et al.	260	501.14			
		Bachovchin et al. Bredehorst et al. Bachovchin et al. Chen et al.	530	331			
			424	94.3	·		
			514	19			
			514	12			
	AJ	5,543,396	08/06/1996	Powers et al.	514	19	
	AK	5,614,379	03/25/1997	MacKellar	435	68.1	
	AL	5,624,894	04/29/1997	Bodor	514	2	
	AM	5,939,560	08/17/1999	Jenkins et al.	548	535	
	AN	6,006,753	12/28/1999	Efendic	128	898	
	AO	5,827,898	10/27/1998	Khandwala et al.	514	734	
	AP	6,201,132 B1	03/13/2001	Jenkins et al.	548	535	04/21/99
	AQ	6,303,661 B1	10/16/2001	Demuth et al.	514	866	04/24/97
	AR	6,500,804 B2	12/31/2002	Demuth et al.	514	19	12/31/02
	AS	6,319,893 B1	11/20/2001	Demuth et al.	514	2	11/20/01
	AT	6,011,155	01/04/2000	Villhauer	544	333	
	AU	6,107,317	08/22/2000	Villhauer	514	365	
	AV	6,110,949	08/29/2000	Villhauer	514	365	
	AW	6,124,305	09/26/2000	Villhauer	514	272	
	AX	6,172,081	01/09/2001	Damon	514	307	
Examiner:				Date:			

INI	FORM PTO-1449 INFORMATION DISCLOSURE STATEMENT			ATTY. DOCKET NO. 20784/4-CON	1	ERIAL NO.	ed
				APPLICANT(S): Demuth et al.			
				FILING DATE: December 11, 2003	b	RT UNIT:	ed
			UNITED STATE	ES PATENT DOCUMENTS			
	AY	5,705,483	01/06/1998	Galloway et al.	514	12	
	AZ	2001/0025023 A1	09/27/2001	Carr	514	2	
	AAA	6,548,481 B1	04/15/2003	Demuth et al.	514	19	11/28/00
	<u>, </u>		FOREIGN P	ATENT DOCUMENTS			
		DOCUMENT NUMBER	DATE	COUNTRY	CLASS	SUB CLASS	TRAN Y/N
	BA	WO 01/62266 A2	08/30/2001	PCT	A61K	38/00	Y
	BB	WO 00/53171	09/14/2000	PCT	A61K	31/155	Y
	ВС	DT 25 42 598 A1	04/22/1976	Germany	C07C	129/16	N
	BD	FR 2 696 740 A1	04/15/1994	France	C07D	207/404	N
	BE	FR 2 085 665	12/31/1971	France	A61K	27/00	N
	BF	WO 97/40832	11/06/1997	PCT	A61K	31/425	Abstract only
	BG	WO 95/15309	06/08/1995	PCT	C07D	207/16	Y
	ВН	JP 4334357	11/20/1992	Japan	C07C	233/57	Abstract only
	BI	WO 93/08259	04/29/1993	PCT	C12N		Y
	ВЈ	WO 95/11689	05/04/1995	PCT	A61K	37/00	Y
	BK	WO 97/45117	12/04/1997	PCT	A61K	31/435	Y
	BL	DE 196 16 486 C2	10/30/1997	Germany	A61K	45/00	Y
	ВМ	WO 95/29691	11/09/1995	PCT	A61K	38/00	Y
	BN	WO 98/22494	05/28/1998	PCT	C07K	5/06	Y
	ВО	WO 00/01849	01/13/2000	PCT	C12Q	1/68	Y
	BP	EP 0 658 568 A1	06/21/1995	EPO	C07K	14/605	Y
	BQ	DD 296 075 A5	11/21/1991	Germany	C07D	295/04	Y
	BR	DD 296 075 A5	11/21/1991	Germany	C07D	295/04	N
	BS	EP 0 708 179 A2	04/24/1996	EPO	C12N	15/16	Y
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	DOCUMENT DATE NUMBER		COUNTRY	CLASS	SUB CLASS	TRAN Y/N	
	ВТ	EP 0 995 440 A1	04/26/2000	EPO	A61K	31/425	N
	BU	WO 91/11457	08/08/1991	PCT	C07K	7/34	Y
	BV	WO 91/16339	10/31/1991	PCT	C07K	5/10	Y
	BW	WO 98/19998	05/14/1998	PCT	C07D	207/00	Y
	BX	WO 91/17767	11/28/1991	PCT	A61K	37/54	Y
	BY	JP 04-288098	10/13/1992	JP	A61K	37/64	(Abstract Only)
	BZ	WO 99/46272 A	09/16/1999	PCT	C07F	9/572	Y
	BAA	DE 299 09 210 U	09/09/1999	Germany	A61K	31/425	N
	BAB	DE 198 26 972 A1	12/23/1999	Germany	A61K	38/05	N
	BAC	WO 01/97808	12/27/2001	PCT	A61K	31/425	Y
	BAD	WO 01/34594 A1	05/17/2001	PCT	C07D	401/06	Y
	BAE	WO 00/10549	03/02/2000	PCT	A61K	31/00	Y
	BAF	WO 01/74299 A2	10/11/2001	PCT	A61K		Y
	BAG	WO 02/20825 A1	03/14/2002	PCT	C12Q	1/00	Y
	BAH	WO 01/89569 A1	11/29/2001	PCT	A61K	45/06	Y
	BAI	WO 99/62914	12/09/1999	PCT	C07F	5/02	Y
	BAJ	WO 99/61431	12/02/1999	PCT	C07D	277/04	Abstract Only-
	BAK	EP 1 130 022 A1	09/15/2001	EPO	C07D	498/09	Y
	BAL	WO 01/32624 A1	05/10/2001	PCT	C07D	217/04	Y
	BAM WO 01/09169 A2 02/08/2001		PCT	C07K	5/062	Y	
	BAN	WO 01/94310 A1	12/13/2001	PCT	C07D	209/12	Y
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	OTHER DOCUMENTS (INCLUDING AU	THOR, TITLE, DATE, PERTINEN	T PAGES, ETC.)		
CA	Campbell, I.W. New Antidiabetic Drugs, ed. efficacy and inadequacy". 3:33-51 (1990).	C.J. Bailey & P.R. Flatt, Smith-Gordon	n, "Sulphonylureas and metformin:		
СВ	CHEMICAL ABSTRACTS, vol. 115. No. 15, 149947q, SCHOEN EKKEHARD ET AL: "D inhibitors on activity of dipeptidyl peptidase I	ipeptidyl peptidase IV in the immune s	system. Effects of specific enzyme		
СС	CHEMICAL ABSTRACTS, vol. 118, no. 25, Hosoda, et al, "Preparation of N-(heterocyclic November 1992 (1992-11-20)				
CD	CHEMICAL ABSTRACTS, vol. 126, no. 2, 1 STOECKEL A. ET AL: "Competitive inhibiti thiazolidides".				
СЕ	Martindale The Extra Pharmacopoeia, 30th Ed	ition, London Pharmaceutical Press, 19	993, Page 1619.		
CF	CF AMASHEH, S., et al., "Electrophysiological analysis of the function of the mammalian renal peptide transporter expressed in Xenopus Laevis oocytes". J. Physiol. 504, 169-174 (1997)				
CG	CG Arai ET AL., "Synthesis of prolyl endopeptidase inhibitors and evaluation of their structure-activity relationships: in vitro inhibition of prolyl endopeptidase from Canine Brain" CHEMICAL AND PHARMACEUTICAL BULLETIN., Bd 41, No. 9, 1993, pages. 1583-1588.				
СН	CH DURINX, C.; et al.; "Reference Values for Plasma Dipepidyl-Pepidase IV activity and their Association with Other Laboratory Parameters". Clin Chem Lab Med 2001, February; 39 (2):155-9, 1 page.				
CI	GOSSRAU, R.; "Cytochemistry of Membrane	e Proteases". Histochem J, 1985, July;	17 (7):737-71, 1 page.		
C1	HAHN, T.; et al.; "Enzyme Histochemical Ev Cultured Villous Explants from Human First T page.				
CK	HEYMANN, E. et al., "Has Dipeptidyl Peptid 1984, January, 2;62 (1):2-10, 1 page.	lase IV an Effect on Blood Pressure and	d Coagulation." Klin Wochenschr,		
CL	CL Lin, J. et al.: "Inhibition of depeptidyl peptidase IV by fluoroolefin-containing n-peptidyl-O-hydroxylamine peptidomimetics" PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES OF USA, Vol. 95, November 1998, pages 14020-14024.				
СМ	KOROM, S., et al "Inhibition of CD26/dipept recipients", <u>Transplantation</u> , Vol. 63, 1495 – 1		ngs cardiac allograft survival in rat		
CN	CN MAGYAR, C.E. et al., "Proximal Tubule Na Transporter Responses are the same during Acute and Chronic Hypertension." Am J. Physiol Renal Physiol, 2000, August; 279 (2):F358-69, 1 page.				
СО	Martindale The Extra Pharmacopoeia, 30th Ed	ition, London Pharmaceutical Press, 19	993, Page 36.		
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•	OTHER DOCUMENTS (INCLUDING A	UTHOR, TITLE, DATE, PERTINEN	T PAGES, ETC.)	
СР	MENTLEIN, R., et al., "Proteolytic processis Pept. 49, 133 -144 (1993)	ng of neuropeptide Y and peptide YY b	y dipeptidyl peptidase IV". Regul.	
CQ	PAPIES, B. et al., "Isoenzyme (Lactate Dehy Activity Changes in Blood Plasma Likely Inc 1991; 33 (3):218-26, 1 page.			
CR	QURESHI. N.U.; et al., "Endogenous Neuro Shock". Regul Pept, 1998, September 25; 7		ring Endotoxic and Hemorrhagic	
CS	TANKA, S., et al., "Suppression of arthritis 19, No. 1 Pages 15-24, (1997)	by the inhibitors of dipeptidyl peptidase	IV". Int. J. Immunopharmacol, Vo	
СТ	The Merck Index, 11th Edition, An Encyclop	edia of Chemicals, Drugs, and Biologic	als, 1989, Page 934	
CU	The Merck Index, 12th Edition, An Encyclop	nedia of Chemicals, Drugs, and Biologicals, 1996, Page 1014.		
CV	Deacon et al., <u>Journal of Clinical Endocrinol</u> Plasma in Vitro Yields and N-Terminally Tr 80(3): 952-957.			
CW	G.G. Duncan, Diseases of Metabolism (Asia	n edition), "Diabetes Mellitus", (1966),	p 951-957.	
CX	Gutniak et al., <u>Diabetes Care</u> , "Subcutaneous Postprandial Glycemia in NIDDM", Septemb		cagon-Like Peptide 1 Abolishes	
CY	Gutniak et al., New England Journal of Med Normal Subjects and Patients With Diabetes		agon-like peptide-1 (7-36) Amide ii	
CZ	H.A. Smith et al., Veterinary Pathology (fou (1972), p 1018-1020.	rth edition), "Diseases and Disorders of	Metabolism: Deficiency Diseases",	
CAA	Hendrick et al., Metabolism – Clinical and E Rats", January 1993, 42(1): 1-6.	xperimental, "Glucagon-like Peptide-I-	(7-37) Suppresses Hyperglycemia in	
CAB	Hoffmann et al., <u>Journal of Chromatography</u> and non-substrate X-X-Pro- oligopeptides as			
CAC	Index Nominum, International Drug Directo	ry 1992/1993, Medpharm Scientific Pub	olishers, pages 728-729.	
CAD	CAD Mannucci et al., <u>Diabetes Care</u> , "Effect of Metformin on Glucagon-Like Peptide 1 (GLP-1) and Nondiabetic Subjects", 24(3): 489-494, March 2001.			
CAE	Nauck et al., <u>Diabetologia</u> , "Normalization of in Type 2 (non-insulin-dependent) diabetic p		glucagon-like peptide 1 (7-36 amid	
CAF	Pauly et al., Regulatory Peptides, "Abstracts Peptides", July 15, 1996, 64(1-3): 148 plus c		onal Symposium on Regulatory	
xaminer:		Date:		

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CAG	Stryer, Biochemistry 3 rd Ed., "Protein Conformation of the Confo	rmation, Dynamics, and Function", 198	8, p 191-193.		
САН	T.J. Kieffer et al., "Degradation of Glucose-I In Vitro and In Vivo by DP IV", Endocrinology		d Truncated Glucagon-Like Peptide 1		
CAI	The Merck Index, An Encyclopedia of Chem	nicals and Drugs, 9th Edition, Merck & 0	Co., Inc., 1976, page 773		
CAJ	C.B. Welch, Medical Management of Non-Ir Association, "Diagnosis and Classification"				
CAK	WETZEL, W., et al., "Effects of the CLIP fra Neuropeptides, 31, 41- 45 (1997)	agment ACTH 20-24 on the duration of	REM sleep episodes".		
CAL	Willms et al., <u>Journal of Clinical Endocrinol</u> Secretion after a Liquid Test Meal: Effects of (Noninsulin-Dependent) Diabetic Patients",	of Exogenous Glucagon-Like Peptide-1			
CAM	CAM Ashworth et al., <u>BIOORG. MED. CHEM. LETT.</u> , "2-Cyanopyrrolidides as Potent, Stable Inhibitors of Dipeptidyl Peptidase IV", (1996), 6(10): 1163-1166.				
CAN	Badia-Elder N.E. et al., <u>Alcoholism Clinical and Experimental Research</u> , "Effects of Neuropeptide Y (NPY) on Ethanolism and Anxiety in High and Low Alcohol Drinking (HAD1/LAD1) Rats", (2000), 24(5): 82A.				
CAO	C.F. Deacon et al., <u>Diabetes</u> , "Both Subcutaneously and Intravenously Administered Glucagon-Like Peptide I Are Rapidly Degraded from the NH ₂ -Terminus in Type II Diabetic Patients and in Healthy Subjects", September 1995, 44: 1126-1131.				
САР	Edwards, J.V. et al., <u>J. Peptide Res.</u> , "Synthe Sequences on Cotton," (1999), 54: 536-543.	sis and Activity of NH ₂ - and COOH-T	erminal Elastase Recognition		
CAQ	Endroczi et al., <u>ACTA PHYSIOL. HUNG.</u> , 'Thymus-Derived Lymphocytes: Effects of In	"Dipeptidyl peptidase IV (DP IV) and Superoxide Dismutase Activity in nhibitory Pepdides and Zn ²⁺ in Vitro", (1990), 75(1): 35-44.			
CAR	CAR Frohman et al., <u>Journal of Clin. Invest.</u> , "Rapid Enzymatic Degradation of Growth Hormone-releasing Hormone by Plasma in Vitro and in Vivo to a Biologically Inactive Product Cleaved at the NH ₂ Terminus", Volume 78, October 198 p 906-913				
CAS	Goodman & Gilman's The Pharmacological	Basis of Therapeutics, Ninth Edition, (1996), p. 1510.			
CAT	Hyun Sook Lee et al., "Cathepsin B Inhibitor	ory Peptides Derived from β-Casein," Peptides 21 (2000) 807-809.			
CAU	Nathan et al., <u>Diabetes Care</u> , "Insulinotropic Subjects", February 1992, 15(2): 270-275.	Action of Glucagonlike Peptide-1-(7-3	7) in Diabetic and Nondiabetic		
CAV	CAV Pauly et al., Metabolism, "Improved Glucose Tolerance in Rats Treated with the Dipeptidyl Peptidase IV (CD26) Inhibitor Ile-Thiazolidide", (1999), 48(3): 385-389.				
CAW	CAW Pschyrembel, Kininisches Wörterbuch 257, Auflage, (1994), 9 pages.				
Examiner:		Date:			

FORM PTO-1449 INFORMATION DISCLOSURE STATEMENT			OOCKET NO. 4/4-CON	SERIAL NO. Not yet assigned	
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	CAX	Snow et al., Advances In Medicinal Chemistr Immunosuppressive Agents", Vol. 3, (1995),		Inhibitors of Dipeptid	lyl Peptidase IV: A New Class of
	CAY	Thorens et al., <u>Diabetes</u> , "Glucagon-Like Pepe NIDDM", (1993), 42:1219-1225.	etide-I and the Co	ontrol of Insulin Secre	tion in the Normal State and in
	CAZ	<u>Vidal</u> , (1993), 69 th Edition, p. 612-613.			, , , , , , , , , , , , , , , , , , ,
	СВА	Wakselman et al., "Inhibition of HIV-1 infect DPP IV activity of CD26", Abstract P 44 of the contract P 44 of the contract P 44 of the contract P			
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